



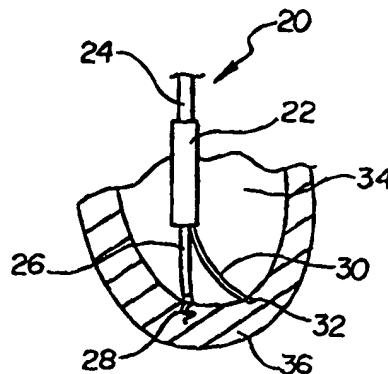
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(54) Title: PERCUTANEOUS MYOCARDIAL REVASCULARIZATION DEVICE AND METHOD

## (57) Abstract

Devices and methods for creating a series of percutaneous myocardial revascularization (PMR) channels in the heart. One method includes forming a pattern of channels in the myocardium leading from healthy tissue to hibernating tissue. Suitable channel patterns include lines and arrays. One method includes anchoring a radiopaque marker (38) to a position in the ventricle wall; then using fluoroscopy repeatedly to guide positioning of a cutting tip (32) in the formation of multiple channels. Another method uses radiopaque material injected into each channel formed, as a marker (38). Yet another method utilizes an anchorable, rotatable cutting probe (30) for channel formation about an anchor. Still another method utilizes a multiple wire radio frequency burning probe, for formation of multiple channels simultaneously. Still another method utilizes liquid nitrogen to cause localized tissue death.



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## PERCUTANEOUS MYOCARDIAL REVASCULARIZATION DEVICE AND METHOD

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### Cross Reference to Related Application

The present application claims the benefit of U.S. Provisional Patent Application No. 60/064,169, filed November 4, 1997.

### Field of the Invention

10        The present application is related to devices and methods for promoting blood circulation to the heart muscle. Specifically, the present invention is related to percutaneous myocardial revascularization (PMR) devices and methods for forming multiple channels in the myocardium.

15

### Background of the Invention

A number of techniques are available for treating cardiovascular disease such as cardiovascular by-pass surgery, coronary angioplasty, laser angioplasty and atherectomy. These techniques are generally applied to by-pass or open lesions in coronary vessels to restore and increase blood flow to the heart muscle.

20        In some patients, the number of lesions are so great, or the location so remote in the patient vasculature that restoring blood flow to the heart muscle is difficult. Percutaneous myocardial revascularization (PMR) has been developed as an alternative to these techniques which are directed at by-passing or removing lesions.

Heart muscle may be classified as healthy, hibernating and "dead". Dead tissue is not dead but is scarred, not contracting, and no longer capable of contracting even if it were supplied adequately with blood. Hibernating tissue is not contracting muscle tissue but is capable of contracting, should it be adequately 5 re-supplied with blood. PMR is performed by boring channels directly into the myocardium of the heart.

PMR was inspired in part by observations that reptilian hearts muscle is supplied primarily by blood perfusing directly from within heart chambers to the heart muscle. This contrasts with the human heart, which is supplied by coronary 10 vessels receiving blood from the aorta. Positive results have been demonstrated in some human patients receiving PMR treatments. These results are believed to be caused in part by blood flowing from within a heart chamber through patent channels formed by PMR to the myocardial tissue. Suitable PMR channels have been burned by laser, cut by mechanical means, and burned by radio frequency 15 current devices. Increased blood flow to the myocardium is also believed to be caused in part by the healing response to wound formation. Specifically, the formation of new blood vessels is believed to occur in response to the newly created wound.

What remains to be provided are improved methods and devices for 20 increasing blood perfusion to the myocardial tissue. What remains to be provided are methods and devices for increasing blood flow to myocardial tissue through controlled formation of channel patterns in the myocardium.

Summary of the Invention

The present invention includes devices and methods for creation of multiple holes in the myocardium of a human heart for percutaneous myocardial revascularization. A pattern of holes is optimally created extending from healthy tissue to hibernating tissue, thereby increasing the supply of blood to hibernating heart muscle tissue. Creating a controlled pattern of channels rather than simply a plurality of channels of unknown location can be accomplished using various methods and devices. Holes can be considered the space left after a volumetric removal of material from the heart wall. Channels have a depth greater than their width and craters have a width greater than their depth.

One method includes marking a first location in the heart muscle wall with a radiopaque marker, then positioning a radiopaque cutting tip relative to the radiopaque marker using fluoroscopy and cutting channels in the myocardium where appropriate. Suitable markers can be secured to the endocardium mechanically with barbs or pigtails or injected into the myocardium. Suitable channel patterns include lines, arrays, and circular clusters of channels.

Another method includes injecting radiopaque material into the newly formed channels, thereby marking the positions of the channels already formed. The radiopaque material should be held in place with polymeric adhesives for the duration of the treatment. The channels formed can be viewed under fluoroscopy using this method. The marker can remain throughout the procedure or only long enough to record the position for mapping.

Yet another method can be accomplished by providing a myocardial channel forming device having an anchoring member, a treatment member with a cutting tip, means for rotating the cutting member about the anchoring member, and means for controlling the radial displacement of the cutting tip from the

5 anchoring member. The anchoring member can be implanted in a heart chamber wall using a pigtail, and the radial and rotational displacement of the cutting tip controlled to sequentially form a circular cluster of channels about the anchoring member. The circular cluster preferably includes both healthy and hibernating tissue areas, which can be mapped using conventional techniques. A variant of

10 this technique utilizes a device having a spline and corresponding star shaft, which restricts the number of possible rotational angles and provide predictable arc rotations around the spline for the treatment member about the anchoring shaft.

Still another method utilizes a bundle of fibers within a sheath as the

15 cutting device. Preferred fibers are formed of Nitinol wire and carry radio frequency current to effect burning channels in the myocardium. Optical fibers carrying laser light for burning are used in another embodiment. The splay of fibers out of the distal end of the sheath can be controlled by controlling the bias of the fibers. The bias of the fibers can be controlled by utilizing shape memory

20 materials, such as Nitinol wire. The splay of fibers can also be controlled by controlling the length of fiber exposed at the distal end, by controlling the retraction of the sheath over the fibers.

A variant device utilizes a magnetically responsive anchoring member, which can be pulled against the heart wall by an external magnetic force. The heart wall can have movement lessened during this procedure and other procedures generally, by inserting a catheter having a magnetically responsive 5 distal region into a coronary artery. Force can be brought to bear upon the heart wall region having the catheter disposed within by applying a magnetic force on the catheter. The applied force can exert a pulling force on the catheter, reducing movement of the beating heart wall in that region.

Another device includes an outer positioning tube having several side 10 channels in the distal region and means for securing the distal region against movement within the heart chamber. One securing means includes a suction orifice near the distal end supplied with vacuum by a vacuum lumen extending the length of the outer tube. Another securing means includes a magnetically responsive portion of the outer tube. The suction orifice can be secured to the 15 heart chamber wall by applying vacuum and the magnetically responsive portion can be forced into the chamber wall by applying an external magnet field. The inner tube can contain an intermediate guide tube and the guide tube can contain an inner PMR cutting wire with a arcuate biased distal region. As the arcuate distal region is moved through the outer tube distal region and over the side 20 channels, the PMR wire distal region can extend through a side channel and to the heart chamber wall. The PMR wire can be moved past undesired side holes by rotating the wire such that the arcuate wire region is oriented away from the side holes.

Another device includes a tube-in-a-tube configuration, having an outer tube disposed about an intermediate tube disposed about an inner PMR cutting probe. The inner PMR probe can be preformed to have a distal region arcuate or angled bias, bent away from the longitudinal axis of the probe. The PMR probe  
5 distal region can extend through a side channel in the distal region of the intermediate tube and is slidable within the intermediate tube, thereby exposing a varying length of distal PMR probe outside of the intermediate tube. The intermediate tube is slidably disposed within the outer tube which has an elongate slot to allow passage of the PMR probe therethrough. Thus, the radial extent or  
10 length of extending PMR probe can be varied by sliding the PMR probe within the intermediate and outer tubes, the longitudinal position of the PMR probe can be varied by sliding the intermediate tube within the outer tube, and the rotational position can be varied by rotating the outer tube from the proximal end. Varying the amount of a preformed, bent PMR probe extending from the intermediate tube  
15 can also change the longitudinal position of the PMR probe distal end.

Another device includes an elongate rod having a distal region secured to an outer collar, such that the outer collar can be pushed and pulled. The outer collar is slidably disposed over an intermediate tube. An inner PMR cutting probe is slidably disposed within the intermediate tube. The inner PMR probe and  
20 intermediate tube together have a distal region arcuate or bent bias or preform, such that distally advancing the outer collar over the intermediate tube straightens out the intermediate tube and proximally retracting the outer collar allows the arcuate bias or bend to be exhibited in the distal region shape of PMR probe and

intermediate tube. The preform can exist in the PMR probe, intermediate tube, or both. The device includes means for anchoring the device to the ventricle wall. Circles or arcs of myocardial channels can be formed by rotating the outer tube, extending the inner PMR probe, and varying the amount of arc to form distal of 5 the outer collar.

Yet another device includes an anchoring member and a positionable cryanoblate treatment tube. The treatment tube can be formed of metal and be either closed or open ended. In use, the device is anchored within a heart chamber and a cryogenic substance such a liquid nitrogen delivered through the 10 tube and to the tube distal end. The liquid nitrogen can cause localized tissue death, bringing about the desired healing response. Still another device includes a plurality of splayed, cryanoblate tubes within a sheath. The tubes can be supplied with liquid nitrogen, which can be delivered through the tube lumens to the tube distal ends so as to cause localized myocardial tissue death at multiple 15 sites substantially simultaneously.

In yet another embodiment, a catheter assembly is provided including a guide wire having a proximal end and a distal end. An expandable member, which may be a wire loop, is disposed at the distal end of the guide wire. The expandable member is moveable between a first position and a second position. 20 In the first position, the member is collapsed to move through a lumen of a guide catheter. In a second position, the expandable member has a transverse diameter, with respect to the length of the guide wire, greater than the transverse diameter of the guide catheter lumen. An elongate catheter having a proximal end and a

distal end is disposed on the guide wire. A therapeutic device is connected to the distal end of the catheter. The therapeutic device can be a needle, hypotube, electrode or abrasive burr to form holes or craters in the myocardium of the patient's heart.

5

Brief Description of the Drawings

Figure 1 is a fragmentary, side, cutaway view of a left ventricle having an anchorable, positionable PMR device within;

Figure 2 is a fragmentary, side view of the PMR device of Figure 1,  
10 showing anchor and treatment members is phantom within a catheter shaft;

Figure 3 is a top view of the PMR catheter and ventricle of Figure 1,  
showing a transverse cross-sectional view of the PMR catheter and a fragmentary  
cross-section and projection of the ventricle wall;

Figure 4 is a fragmentary, perspective view of a multiple-tip PMR  
15 treatment device according to the present invention;

Figure 5 is an end view of the multiple-tip PMR treatment device of  
Figure 4;

Figure 6 is a fragmentary, side, cutaway view of a left ventricle having a  
magnetically anchorable, positionable PMR device within;

20 Figure 7 is cutaway, perspective view of a heart having a magnetically  
positionable PMR cutting tip within the left ventricle wall;

Figure 8 is a perspective view of a heart having a magnetic, heart wall  
stabilizing catheter disposed within the left coronary artery, shown in phantom;

Figure 9 is a perspective view of a multiple channel positioning device for forming multiple myocardial channels in a ventricle wall, having distal anchoring means and containing a guide catheter containing a PMR cutting wire, both drawn in phantom;

5       Figure 10 is a fragmentary, perspective view of a device related to the device of Figure 9, illustrated without distal anchoring means, better illustrating a shape member within the device;

10      Figure 11 is a perspective view of a tube-in-a-tube positioning device for positioning a PMR cutting probe, having an outer tube containing an inner tube containing a PMR cutting probe;

Figure 12 is a fragmentary, perspective view of a section through the PMR probe of Figure 11, better illustrating the shape member;

15      Figure 13 is a perspective view of an extendable collar device for positioning a PMR probe, having a slidable collar over an intermediate tube over a PMR cutting probe;

Figure 14 is a fragmentary, side, cutaway view of a left ventricle having an anchorable, positionable cryanoblative PMR device within;

Figure 15 is a fragmentary, perspective view of a multiple-tip cryanoblative PMR treatment device according to the present invention;

20      Figure 16 is a perspective view of yet another embodiment of the device in accordance with the present invention;

Figure 17 is a view of the device of Figure 16 in use;

Figure 18 is an alternate embodiment of the device of Figure 16;

Figure 19 is an alternate embodiment of the device of Figure 16; and

Figure 20 is an alternate embodiment of the device of Figure 16.

Detailed Description of the Preferred Embodiments

5       Figure 1 illustrates an anchorable percutaneous myocardial revascularization (PMR) treatment catheter 20 disposed within a left ventricle 34. PMR catheter 20 includes an inner star shaft 24 disposed within an outer catheter shaft 22, an anchoring shaft 26 disposed within star shaft 24, and a treatment shaft or probe 30 disposed within catheter shaft 22. Catheter shaft 22 has been cut 10 away proximally in Figure 1, illustrating inner star shaft 24 within. Anchoring shaft 26 has an anchor 28 disposed at the distal end. In a preferred embodiment, anchor 28 has a pigtail or corkscrew configuration, capable of reversibly securing itself to the ventricular wall through rotation of anchoring shaft 26. One embodiment anchor includes a distal barb, capable of securing itself to the 15 ventricular wall through translation of anchoring shaft 26, not requiring shaft rotation for anchoring. In another embodiment, anchoring shaft 26 includes a vacuum lumen therethrough terminating in a distal orifice or suction tip (not shown). Treatment shaft 30 has a distal cutting tip 32, shown embedded within a section of a left ventricular wall 36. The term "cutting" as used herein includes 20 penetrating and channel forming by other means.

Referring now to Figure 2, PMR catheter 20 is illustrated in more detail. Anchor shaft 26, extending through outer catheter shaft 22, includes a distal radiopaque marker 38. Treatment shaft 30, extending through catheter tube 22,

preferably includes an arcuate, distal region 33 and a distal radiopaque marker 40.

Radiopaque markers 38 and 40 can aid in determining the positions of the anchoring and treatment shafts under fluoroscopy. Suitable radiopaque materials are well known to those skilled in the art, including barium, bismuth, tungsten and

5 platinum. Referring now to Figure 3, PMR catheter 20 is illustrated in a top, cross-sectional view taken through the catheter. In a preferred embodiment, anchoring shaft 26 is contained within an anchor shaft lumen 27. Anchor shaft lumen 27 is preferably slidably disposed within an inner shaft such as star shaft 24. Inner shaft 24 preferably has a star shape and is disposed within a star lumen  
10 25 having internal splines corresponding to the vertices of star shaft 24. Treatment shaft 30 is preferably slidably disposed within a treatment shaft lumen 31 within the wall of PMR outer shaft 22. As illustrated, treatment shaft 30 cutting end 32 has formed several channels 42 in the myocardium of ventricular wall 36.

15 The use of PMR device 20 may now be discussed, with reference to Figures 1, 2 and 3. Several degrees of freedom of movement of cutting tip 32 are possible with the present invention. Treatment shaft distal region 33 is preferably biased to assume a more radially extended position when unconstrained by lumen  
31. Cutting tip 32 may be seen to have a radial distance "R" from anchoring shaft  
20 26, as indicated in Figure 2. Holding the axial displacement of anchoring shaft 26 and treatment shaft 30 fixed while distally, axially sliding catheter outer shaft 22 over both shafts 26 and 30 causes more of treatment shaft distal region 33 to be drawn into outer shaft 22, thereby decreasing the radial distance R of cutting tip

32 from anchoring shaft 26. Thus, by proximally fixing the longitudinal positions of anchoring shaft 26 and treatment shaft 30, and sliding outer shaft 26 over a range of motion, a series of channels along a line extending radially outward from anchoring shaft 26 can be created. It will be recognized that, to the extent the 5 inner ventricular wall does not match the arcuate shape of treatment shaft distal region 33, it may be necessary to adjust the longitudinal displacement of treatment shaft 30 within outer shaft 22 as well, to enable cutting tip 32 to reach the endocardium.

In addition to cutting a series of channels radially outward from anchoring 10 shaft 26, cutting tip 32 can also describe an arc about treatment shaft lumen 31, best visualized with reference to Figure 3. By rotating treatment shaft 30 within lumen 31, cutting tip 32 can sweep through an arc, cutting a regular series of channels into the myocardium. By varying radial distance R and the rotation of treatment shaft 30, a regular series of arcs of channels can be formed, with the 15 arcs having increasing radial distance from outer shaft 22.

Outer shaft 22 can also be rotated relative to anchoring shaft 26, thereby enabling the cutting of a regular series of channels in a circle about anchoring shaft 26. In a preferred embodiment, an intermediate star shaft such as shaft 24 is disposed between anchoring shaft 26 and outer shaft 22. Star shaft 24 can serve 20 to restrict the rotational positions possible for outer shaft 22 relative to inner, anchoring shaft 26. Outer shaft 22 having internal splines, is not freely rotatable about the vertices of star shaft 24. In order for outer shaft 22 and carried treatment shaft 30, to be rotated about anchor shaft 26, star shaft 24 can be star

shaped only is a limited distal region, and outer shaft 22 only splined in a limited distal region. In a preferred embodiment, star shaft 24 and outer shaft 22, at a location proximal of the cross section of Figure 3, have smooth outer and inner surfaces, respectively. The smooth surfaces allow star shaft 24 to be rotated 5 within outer shaft 22 when star shaft 24 has been retracted proximally into the smooth region. After rotation, star shaft 24 can be advanced distally, sliding within a spline of outer shaft 22. The rotation of outer shaft 22 can thus be restricted when desired and enabled when desired. When enabled, rotation of shaft 22 can thus be restricted to a discrete set of rotational angles. Another 10 embodiment of the invention dispenses with intermediate, start shaft 24, allowing outer shaft 22 to rotate directly about inner, anchoring shaft 26. In this embodiment, the rotation of outer shaft 22 about anchoring shaft 26 is not restricted to a set of discrete rotational angles.

Cutting tip 32 can form a substantially regular pattern of channels. 15 Cutting tip 32 preferably is formed of a wire such as Nitinol or elgiloy or stainless steel, and is capable of delivering the radio frequency current used for cutting channels in the myocardium. A suitable device for radio frequency cutting is described in co-pending U.S. Patent Application Serial No. 08/810,830, filed March 6, 1997, entitled RADIOFREQUENCY TRANSMYOCARDIAL 20 REVASCULARIZATION APPARATUS AND METHOD. By restricting the movement of cutting tip 32 to movements relative to anchor tip 28, a more regular pattern of channels can be formed, even with limited fluoroscopic feedback,

relative to the pattern formed by a cutting tip operating independent of the anchoring tip.

Referring now to Figure 4, a multi-fiber treatment probe 50 is illustrated. Treatment probe 50 includes a plurality of wires or optical fibers 54, having distal cutting tips 56, and enclosed within a sheath 52. Figure 5 illustrates an end view of multi-fiber probe 50, showing distal cutting tips 56 in the pattern they would have approaching the myocardium. Probe 50 allows a pattern of channels to be formed in the myocardium at the same time, not requiring repeated re-positioning of a single cutting tip such as cutting tip 32 of Figure 2. Wires 54 are preferably formed of Nitinol wire. Use of a bundle of fibers including metal wires or optical fibers allows use of RF or laser cutting means, respectively. RF and laser cutting allows use of fibers relatively close together, as illustrated in Figure 5. Mechanical cutting tips, such as those using rotating cutting blades, can require more space between cutting tips, not allowing the dense coverage of Figure 5. In one embodiment, the cutting tips have an outside diameter "D" and an average inter-strand distance "I", as illustrated in Figure 5, where I is about 2 to 3 times the value of D. The pattern of cutting tips can be controlled by utilizing radially outwardly biased cutting tips, which splay outward as illustrated in Figure 4. The amount of splay is controlled in one embodiment by allowing the enclosing sheath 20 to retract, allowing the cutting tips to splay further outward. Sheath 52 can prevent uncontrolled flopping of distal cutting tips 56, which can present a problem when large inter-strand distances are required, as with some mechanical cutting tips. The coverage of the cutting tips in Figure 5 allows creation of a

complete pattern of channels in the myocardium without requiring repositioning of the cutting tips.

In use, probe 50 can be positioned near the ventricle wall region to be revascularized, and RF current delivered through distal cutting tips 56. The 5 resulting myocardial channels can be formed substantially at the same time, and a similar pattern delivered to an adjacent ventricular wall area soon thereafter.

In another embodiment of the invention, not requiring illustration, a radiopaque marker can be delivered and secured to a position in the ventricular wall. Suitable radiopaque materials include barium, bismuth, tungsten and 10 platinum. Markers believed suitable include metal markers having barbs or pigtails to securely engage the ventricle wall. Other markers, such as radiopaque gels injected into the ventricular wall, are suitable provided they stay in place for the length of the procedure. Such markers are preferably injected from within the ventricle utilizing a catheter. A preferred method utilizes the cutting tip to first 15 plant or inject a marker, followed by the cutting of a series of channels in the myocardium. By utilizing a radiopaque distal cutting tip and a fixed, implanted radiopaque marker, the relative positions of the two can be viewed fluoroscopically and adjusted fluoroscopically, thereby allowing formation of a controlled pattern of channels. The radiopaque marker provides a reference point 20 for forming a pattern of channels in the myocardium.

In another embodiment of the invention, the cutting tip injects radiopaque material in conjunction with the cutting of a channel. In this embodiment, as each channel is formed, a radiopaque marker is left, creating a pattern of radiopaque

markers viewable fluoroscopically. The pattern of channels formed in the myocardium are thus immediately viewable, giving feedback to the treating physician as to the progress and scope of the pattern of channels. Suitable materials for injection into the myocardium are preferably biodegradable or  
5 absorbable into the body soon after the procedure, allowing the myocardial channels to be perfused with blood. A device suitable for cutting and injection of material is described in copending U.S. Patent application Serial No. 08/812,425, filed March 16, 1997, entitled TRANSMYOCARDIAL CATHETER AND METHOD, herein incorporated by reference.

10 Referring now to Figure 6, left ventricle 34 having a magnetically anchorable, positionable PMR device 86 device within is illustrated. PMR device 86 is similar in some respects to PMR device 20 illustrated in Figure 1, with device 86 differing primarily at the distal end of anchoring shaft 26. Anchoring shaft 26 has a magnetically responsive portion 80 at the anchoring shaft distal end. "Magnetically responsive" as used herein refers to a material capable of being attracted or repelled by a magnet. Magnetically responsive portion 80 can be used in conjunction with external magnets to position anchoring shaft 26 against the ventricle wall. External magnets such as magnet 84 can be disposed external to the body, positioned to direct the distal end of anchoring shaft 26 into  
15 the center of a target area in the heart. In one embodiment, the external magnets are rare earth magnets. In another embodiment, the external magnets are superconducting magnets. In a preferred embodiment, several magnets 84 are used to direct anchoring shaft 26 into the heart wall.  
20

In use, magnets 84 can be used in conjunction with axially moving anchoring shaft 26 to plant anchoring shaft 26 in the desired location. Pairs of magnets in all three dimensions may not be required as the goal is to pull the anchoring shaft against a ventricle wall, not necessarily to suspend it in place

5 using the magnets. The magnets, in conjunction with a radiopaque anchoring shaft tip and fluoroscopy, can be used to guide the anchoring shaft into position and maintain position during treatment. In the embodiment illustrated, an anchoring spike 82 lies at the distal end of anchoring shaft 26. Anchoring spike 82, drawn larger in Figure 6 than in the preferred embodiment, is used to stabilize

10 the position of the anchoring shaft distal end once the desired position has been reached. Another embodiment terminates anchoring shaft 26 without any spike, rather ending with magnet 80. Still another embodiment terminates anchoring shaft 26 with an orifice, such as a suction tip, in communication with a vacuum lumen within shaft 26, allowing anchoring shaft 26 to be held in place by

15 applying vacuum to the vacuum lumen and orifice, thereby securing the distal tip of shaft 26 with vacuum pulling against the heart chamber wall.

Referring now to Figure 7, a heart 35 having a PMR catheter 90 disposed within. PMR catheter 90 includes a shaft 92, illustrated extending through the aorta and into left ventricle 34. A magnetically response distal portion 94 is located near a distal cutting tip 96 on PMR catheter 90. As illustrated, cutting tip 96 has been guided into left ventricular wall 36 and has cut a channel in the wall. External magnets 84 can be used to position cutting tip 96 into the desired position with the aid of fluoroscopy. Distal portion 94 is preferably radiopaque,

to aid in guiding cutting tip 96 into position. As PMR catheter shaft 92 provides some degree of support to cutting tip 96, and as the primary goal is to pull cutting tip 96 into the ventricular wall, pairs of magnets in all three dimensions may not be required. External magnets 84 serve to position cutting tip 96, and, with the 5 assistance of catheter shaft 92, can serve to pull cutting tip 96 into the ventricular wall.

Referring now to Figure 8, a magnetically responsive catheter 100 is illustrated, disposed within heart 35, being extended through aorta 102 into a left coronary artery 104. Catheter 100 includes a magnetically responsive distal 10 region 106, which can be attracted by external magnets 84. Catheter 100 can be used in conjunction with external magnets to stabilize regions of the heart, lessening the amount of wall movement due to the beat of the heart.

In use, magnetically responsive catheter 100 can be advanced with aid of fluoroscopy through the aorta and into a coronary artery. Catheter distal region 15 106 preferably includes radiopaque materials to aid positioning under fluoroscopy. Once in position, distal region 106 is effectively located in the heart wall. When stabilization is desired, external magnets such as magnet 84 can be positioned near catheter distal region 106. By exerting a strong pull on distal region 106, the movement of the heart wall in the vicinity of catheter distal region 20 106 can be lessened.

Stabilization can be used during intravascular PMR procedures, minimally invasive PMR procedures, and heart procedures generally. When used during PMR procedures, the stabilization can serve to lessen heart wall movement in the

area being cut. When used during other medical procedures, the stabilization can serve to minimize heart wall movement in areas being operated on or otherwise treated. When used during intravascular PMR procedures, a second, PMR catheter should be provided.

5 Referring now to figure 9, a multiple-channeled PMR positioning device or guiding tube 120 is illustrated. Positioning tube 120 includes a distal end 122, a distal region 124, a proximal end 126, a plurality of channels 138 within distal region 124, and a lumen 128 therethrough. A distal anchoring means 130 is preferably located distal of distal region 124 and can serve to fix the position of  
10 distal end 122 to the wall of the left ventricle or other heart chamber. In one embodiment, anchoring means 130 includes an orifice or suction tip in communication with a vacuum lumen 148, such that anchoring means 130 can be held in place against a heart chamber wall once positioned near the wall. In another embodiment, anchoring means 130 includes a magnetically responsive  
15 material such that an externally applied magnetic field can force anchoring means 130 into a heart chamber wall. In this magnetically responsive embodiment, anchoring means 130 can be similar to distal portion 94 illustrated in Figure 7. In another embodiment, tube distal region 124 is magnetically responsive and can be similar to magnetically responsive region 106 illustrated in Figure 8. Distal tip  
20 122 is preferably formed of soft,atraumatic material and distal region 124 formed of sufficiently pliable material so as to allow distal region 124 to conform to a ventricle wall.

Disposed within positioning tube 120 is a guide catheter 142 extending from positioning tube proximal end 126 to distal region 124. Disposed within guide catheter 142 is a PMR cutting wire 132, proximally electrically connected to an RF energy source 136 and terminating distally in a cutting tip 33. PMR wire 5 132 includes a distal arcuate or bent region 144 proximate distal cutting tip 33. Arcuate region 144 can be bent or arced so as to have a preformed shape or bias to extend laterally away from the longitudinal axis of the PMR wire. In one embodiment, PMR wire lies within positioning tube 120 directly, without a guide catheter. In a preferred embodiment, a guide catheter such as guide catheter 142 10 is disposed about the PMR wire. PMR wire 132 is slidably disposed within guide catheter 142 and can be rotated by applying torque to the proximal end.

In use, positioning tube 120 can be preloaded with guide catheter 142 containing PMR wire 132. PMR wire 132 can be retracted such that arcuate region 144 is retracted either to a position proximal of channels 138 or within 15 positioning tube distal region 124 but retracted within guide catheter 142. In this retracted position, PMR wire arcuate region 144 does not extent from channels 138. With PMR wire retracted, positioning tube 120 can be advanced through the vasculature into a heart chamber such as the left ventricle. Positioning tube distal end 122 can be advanced down into the ventricle and up a ventricular wall. With 20 distal end 122 in a desired position, anchoring means 130 can be used to anchor distal end 122 to the ventricular wall. In embodiments where anchoring means 130 is magnetically responsive or where positioning tube distal region 124 is magnetically responsive, an external magnetic force can be applied to pull or push

anchoring means 130 and distal region 124 into the wall. In embodiments where anchoring means 130 is a suction tip, vacuum can be applied to the vacuum lumen in communication with the suction tip.

With positioning tube distal region 124 in place, guide catheter 142 containing PMR wire 132 can be advanced to push tube distal end 122 and PMR wire arcuate region 144 distally out of guide catheter 142. PMR wire 132 can be rotated such that cutting tip 33 is oriented toward channels 138, and guide catheter 142 and PMR wire 132 retracted together until cutting tip 33 can be pushed out of channel 138. Cutting tip 33 can be advanced through channel 138 and a channel 10 cut into the myocardium. In a preferred embodiment, PMR wire 132 has a depth stop 146 proximal of arcuate region 144 that limits the length of wire passed through channels 138, such that the depth of a PMR formed myocardial channel is limited. After myocardial channel formation, PMR wire 132 can be retracted through the channel and the next, more proximal channel entered. In a preferred 15 embodiment, arcuate region 144 is radiopaque and a series of radiopaque marker bands separate channels 138 to aid in positioning cutting tip 33. In one embodiment, PMR wire 132 can be rotated to cut more than one myocardial channel per positioning tube channel. In this manner, a series of myocardial channels in a regular pattern can be formed over the length of positioning tube 20 distal region 124.

Referring now to Figure 10, another embodiment positioning tube 160 is illustrated. Positioning tube 160 has a shape member 164 which can assist in forming the U-shape of tube 160 illustrated in Figure 10. In one embodiment,

shape member 164 is formed of a shape memory material such as Nitinol and embedded within the wall of tube 160 to impart a shape to the tube once tube 160 is within a ventricle and is no longer as restrained as when disposed within a blood vessel or guide catheter. In another embodiment, shape member 164 is a

5 pull wire slidably disposed within a lumen within tube 160 and fixedly attached to a distal portion of the tube as indicated at 166. In this embodiment, shape member 164 can be pushed and pulled from a proximal location outside of the patient's body so as to assist in imparting a shape to tube distal region 124. In

10 Figure 10, the distal most portion of tube 160, including anchoring means 130, has been omitted from the drawing to more clearly illustrate the distal termination of shape member 164. From inspection of Figure 10, it may be seen that, by rotating positioning tube 160 to different anchoring positions, and by advancing PMR wire 132 to various tube channels, a large expanse of ventricular wall can be covered and have myocardial channels formed therein.

15 Referring now to Figure 11, a tube-in-a-tube embodiment positioning device 180 is illustrated. Positioning device 180 includes an inner PMR cutting probe 182 slidably disposed within an intermediate tube 184 which is slidably disposed within an outer tube 186. PMR probe 182 has a cutting tip 188 and preferably has radiopaque marker bands 190. Marker bands 190 aid in

20 positioning the PMR probe under fluoroscopy. PMR probe 182 is preferably preformed to have an arcuate or bent distal region 192.

Intermediate tube 184 has a channel 194 formed through the tube wall sufficiently large to allow passage of PMR probe 182. In a preferred

embodiment, channel 194 is formed in a side tube wall in a distal portion of intermediate tube 184, as illustrated in Figure 11. Outer tube 186 has an anchoring tip 200 and a slot 196, with slot 196 illustrated extending along the longitudinal axis of the outer tube. Slot 196 is sufficiently wide to allow passage 5 of PMR probe 182 therethrough. In one embodiment, anchoring tip 200 is formed of a soft material and held in place by axial force directed along the longitudinal axis of device 180. In another embodiment, anchoring tip 200 contains a magnetically responsive material and is held in place at least partially by externally applied magnetic forces. Referring now to Figure 12, a section of PMR 10 probe 182 is further illustrated, showing one structure for imparting a preformed arc or bend to the probe. PMR probe 182 can include a tube wall 199 having a preform wire 198 embedded therein. Preform wire 198 is preferably formed of a shape memory material such as Nitinol, such that the arcuate or bent shape is reformed upon exit from the constraint of intermediate tube 184.

15 Referring again to Figure 11, the wide range of motion possible for cutting tip 188 may be discussed. The radial extent of cutting tip 188, the distance from the center longitudinal axis of outer tube 186, can be varied by extending PMR probe 182, thereby forcing a longer extent of exposed probe through intermediate tube channel 194 and through outer tube slot 196. As PMR probe 182 has arcuate 20 region 192 in a preferred embodiment, extending PMR probe also changes the longitudinal position of the cutting tip as more arc is exposed. Sliding intermediate tube 184 within outer tube 186 also changes the longitudinal position of cutting tip 188. Cutting tip 188 is illustrated at a first position A in Figure 11, a

second, more distal position B, and a third, still more distal position C, as intermediate tube 184 is advanced distally within outer tube 186. Finally, outer tube 186 can be rotated about its center, longitudinal axis, thereby extending the range of coverage of cutting tip 188.

5 In use, PMR positioning device 180 can be advanced into the left ventricle and anchoring tip 200 forced against some portion of the vermicular wall. Intermediate tube 184 can be slid within outer tube 186 to a desired position. Inner PMR probe 182 can be advanced out of channel 194 until the desired length of PMR probe is exposed. A desired position of cutting tip 188 can be reached by

10 10 adjusting the length of PMR probe 182 exposed, the length of intermediate tube 184 advanced into outer tube 186, and the rotation of outer tube 186. In one method, a series of arcs of myocardial channels are formed substantially transverse to the longitudinal axis of positioning device 180. In this method, outer tube 186 is rotated such that cutting tip 188 describes an arc. As each arc is

15 completed, intermediate tube 184 is slid relative to outer tube 186 and a new arc of channels is burned into the ventricular wall.

Referring now to Figure 13, an extendable collar embodiment PMR positioning device 220 is illustrated. Device 220 includes inner PMR probe 182 disposed within an intermediate tube or sleeve 222 which is slidably disposed within an outer collar 224. Intermediate sleeve 222 includes a distal end 240 and has a lumen 242 extending therethrough. Inner PMR probe 182 is preferably slideable within intermediate sleeve 222. Device 220 includes an elongate rod 226 having a distal region 228 secured to outer collar 224. In a one embodiment,

elongate rod 226 is capable of both pulling and pushing outer collar 224 over intermediate sleeve 222. An elongate anchoring member 230 includes a proximal region 236, a distal end 234, a distal anchoring means such as pigtail 234, and can be slidably and rotatably secured to outer collar 224.

5 In one embodiment, elongate rod 226 and anchoring member 230 are both slidably disposed in a dual lumen tube 227 substantially coextensive with intermediate tube 222. Dual lumen tube 227 can terminate the lumen containing elongate rod 226 in a skived portion 229, continuing the tube as a single lumen portion 231. Single lumen portion 231 allows elongate rod 226 to freely travel  
10 with outer collar 224. Outer collar 224 preferably is slidably disposed over single lumen portion 231.

Intermediate sleeve 238 and inner PMR probe 182 together have an arcuate or bent bias or preform, as illustrated at 238. In one embodiment, intermediate sleeve 222 has a preformed shape which can be imparted with an  
15 embedded shape wire as illustrated by wire 189 in Figure 12. In another embodiment, PMR probe 182 has an arcuate bias sufficiently strong to impart a distal bend to both intermediate sleeve 238 and PMR probe 182 when outer collar 224 is retracted. PMR probe 182 can include Nitinol or other shape memory material to impart this arcuate bias. In yet another embodiment, both inner PMR  
20 probe and intermediate sleeve 238 have a preformed arcuate or bent distal shape.

In one embodiment, intermediate tube 222 can be rotated within outer collar 224. In another embodiment, intermediate tube is restricted in rotation corresponding ridges and grooves between outer collar 224 and intermediate tube

222. In one embodiment, outer collar has internal ridges fitting within external grooves in a region of intermediate tube 222. Restricting the rotation of intermediate tube 222 within collar 224 can aid in causing rotation about anchoring member 230 rather than about the center of outer collar 224.

5       In use, outer collar 224 can be extended distally over intermediate sleeve 222, such that collar 224 is proximate intermediate sleeve distal end 240. Inner PMR probe 182 can be preloaded within intermediate sleeve 222. With outer collar 224 distally extended, arcuate region 238 is substantially restrained and straightened. Device 220 can be advanced within the vasculature and into a heart 10 chamber such as the left ventricle. Elongate anchoring member 230 can be advanced distally and rotated, thereby rotating pigtail 232 into the ventricle wall and securing anchoring member 230. With intermediate sleeve 222 and PMR cutting tip 188 positioned as indicated at "E" in Figure 13, the extent of PMR probe exposed can be adjusted by axially sliding PMR probe 182 within 15 intermediate sleeve 222. The extent of intermediate sleeve extending distally beyond collar 224 can be adjusted in some embodiments by advancing or retracting sleeve 222 within collar 224. With PMR cutting tip 188 in position, intermediate sleeve 222 can be rotated about anchor member 230 and a circular pattern of myocardial channels can be burned about the pigtail. In a variant 20 method, possible in devices allowing rotation of intermediate sleeve 222 within outer collar 224, intermediate sleeve 222 can be rotated about the center axis of outer collar 224. With one circle completed, outer collar 224 can be retracted, allowing more of the preformed shape of sleeve 22 and PMR probe 182 to appear,

as illustrated, for example, at "D" in Figure 13. As collar 224 is retracted, PMR probe 182 can be advanced to describe circular paths of increasing radius over the inner ventricle walls. In this way, a series of circular paths of myocardial channels about the anchoring point can be formed in the ventricle walls. In one embodiment, elongate member 226 is capable of only retracting collar 224, which, once retracted within the ventricle, cannot be advanced within the ventricle. In another embodiment, elongate member 226 is capable of both advancing and retracting collar 224 over intermediate sleeve 222. With the formation of myocardial channels complete, anchoring member 226 can be rotated opposite the initial rotation, thereby releasing pigtail 232 from the ventricle wall.

Figure 14 illustrates an anchorable, cryanoblative PMR treatment catheter 320 disposed within a left ventricle 34. The term "cryanoblative", as used herein, refers to the delivery of cold sufficient to cause tissue death. Similarly numbered elements are discussed with respect to Figure 1. Cryanoblative catheter 320 includes an inner star shaft 24 disposed within an outer catheter shaft 22, an anchoring shaft 26 disposed within star shaft 24, and a cryanoblative treatment tube 330 disposed within catheter shaft 22. Cryanoblative treatment tube 330 is preferably formed of metal and can include a distal cryanoblative tip 332 and a lumen through which a cold substance, such as liquid nitrogen, is delivered.

In one embodiment, distal cryanoblative tip 332 includes a distal orifice in communication with the treatment shaft lumen, such that liquid nitrogen can be delivered through the orifice and to the heart chamber wall. In another

embodiment, tube 330 is close ended and initially under vacuum, allowing liquid nitrogen to be delivered to the tube distal region, causing the tube to become very cold without allowing liquid nitrogen to enter the myocardium. The cryanoblate tip can be inserted into the heart chamber wall, penetrating the wall, and into the 5 myocardium prior to delivery of liquid nitrogen. The delivery of liquid nitrogen to the heart chamber wall can cause localized tissue death, bringing about the same healing response as laser and radio-frequency current PMR.

Referring now to Figure 15, a multi-tube, cryanoblate treatment probe 350 is illustrated. Treatment probe 350 includes a plurality of cryanoblate tubes 10 354, having distal cryanoblate cutting tips 356, and enclosed within a sheath 52. In one embodiment, tubes 354 are feed from a common supply within sheath 52, such that tubes 354 have a short length, with most of the length lying distal of the sheath. Probe 350 allows a pattern of channels to be formed in the myocardium at the same time, not requiring repeated re-positioning of a single cutting tip such as 15 cutting tip 332 of Figure 14. The pattern of cutting tips can be controlled by utilizing radially outwardly biased cutting tips, which splay outward as illustrated in Figure 15. The amount of splay is controlled in one embodiment by allowing the enclosing sheath to retract, allowing the cutting tips to splay further outward. Sheath 52 can prevent uncontrolled flopping of distal cutting tips 356, which can 20 present a problem when large inter-strand distances are required, as with some mechanical cutting tips.

The coverage of the cutting tips in Figure 15 allows creation of a complete pattern of channels in the myocardium without requiring repositioning of the

cutting tips. The resulting myocardial channels can be formed substantially at the same time, and a similar pattern delivered to an adjacent ventricular wall area soon thereafter. As discussed with respect to Figure 14, cryanoblative tubes 354 can be formed of metal and be either closed or open ended.

5 In a variation of the methods previously described, a radiopaque contrast media is used to determine the depth of channels formed in the myocardium. The contrast medium is injected or "puffed" into or near the channel formed in the myocardium. The heart can be visualized under fluoroscopy to determine the depth of the channel formed thus far. After visualization, the channel can be  
10 further deepened. The cycle of channel formation, contrast medium puffing, and fluoroscopic visualization can be repeated until the channel has the desired depth.

Contrast medium could be injected using a lumen such as the lumen of guide catheter 142 of Figure 9. A lumen such as the lumen in tube 330 of Figure 14 could also be used to deliver contrast medium. The lumens previously  
15 discussed with respect to injecting liquid nitrogen could be used to deliver contrast medium.

In addition to using the device as described herein above to form channels in the myocardium, the device could be used to form craters in the myocardium. That is, to form a wound in the myocardium having a width greater than its depth.  
20 The crater can be formed by controlling the depth of insertion of, for example, a radiofrequency device and/or controlling the power delivered to the distal tip of the device such that a crater is formed. Those skilled in the art can also appreciate that mechanical devices, laser devices or the like could be used to form craters.

In use, the above methods and devices can be used to form a pattern of channels leading from healthy myocardial tissue to hibernating tissue. This can operate by multiple mechanisms to supply hibernating tissue with an increased blood supply. First, channels in the myocardium can perfuse tissue directly from

5 the ventricle, through the patent channel formed by the cutting tip. Second, the channels formed by the cutting tip can become newly vascularized by operation of a healing response to the channel injury. The new blood vessels thereby increase further the supply of hibernating tissue by ventricular blood. Third, the series of

10 newly formed vessels caused by the healing response can form interconnections or anastomoses between the series of injured areas, forming a network of blood vessels, which, by connecting with healthy area vessels, can be supplied by blood originating from coronary arteries in addition to blood supplied directly by the

15 ventricle.

Figure 16 shows yet another embodiment of the present invention in the

15 form of catheter assembly 400. Here only the distal end of catheter assembly 400 is shown disposed within the left ventricle of a patient's heart. Those skilled in the art will appreciate the various configurations possible for the proximal end of the catheter in view of the description of the distal end which follows. Catheter assembly 400 includes an elongate guide wire 402 having a distal end and a

20 proximal end. A collapsible loop 404 is hingably connected to the distal end of guide wire 402. A retraction member 406 is hingably connected to loop 404 opposite the connection to guide wire 402. Therapeutic catheter 408, which has a lumen extending therethrough, is shown advanced over guide wire 402. Catheter

408 has a distal end and a proximal end, and proximate the distal end of catheter 408 is a therapeutic member 410. Therapeutic member 410 can be an elongate electrode having a ball tip. In a preferred embodiment, a conductor extends through catheter 408 to deliver RF energy to electrode 410. Electrode 410 can be 5 hingably connected to catheter 408 such that catheter assembly 400 can be advanced through a guide catheter 412.

The materials to be used, and the methods of fabrication, to make catheter assembly 400 will be known to one skilled in the art in view of the uses to which catheter assembly 400 are put. As shown in Figure 16, loop 404 is disposed in a 10 first collapsed position A. In collapsed position A, loop 404 is advancable to left ventricle 34 of heart 35 by a percutaneous route through the aorta. Figure 17 shows loop 404 in a second position B deployed within left ventricle 34. When loop 404 is in second position B, a portion of guide wire 402 lies near and approximately parallel to left ventricle wall 36, while a portion of loop 404 abuts 15 the opposite wall. In this position, catheter 408 can be advanced as shown by the arrow along guide wire 402. As catheter 408 is advanced, electrode 410 can be energized repeatedly to form holes or channels 442 in wall 36. A further series of holes 442 can be formed by rotating wire 402 and loop 404 as shown by the arrows adjacent loop 404.

20 In order to move loop 404 between first position A and second position B, guide wire 402 should be relatively rigid in comparison to loop 404 and actuator member 406. With that configuration, actuator 406 can be pulled proximally to

move loop 404 from second position B to first position A. In turn, actuator member 406 can be moved distally to deploy loop 404.

Figure 18 is a view of the distal end of catheter 408 showing an alternate therapeutic device. In particular, a hypodermic needle 414 is shown extending 5 from distal end 408. Hypodermic needle 414 is preferably hingable connected to catheter 408 such that it can be advanced and withdrawn through a guide catheter. If catheter 408 includes an infusion lumen, contrast media, growth factor or other drug can be delivered to wall 36 through needle 414.

Figure 19 is a view of the distal end of catheter 408 showing yet another 10 therapeutic device disposed thereon. In particular, an electrode 416 is shown which has a length greater than the distance which it extends transversely from catheter 408. Such an electrode can be used to form a crater 444 having a width greater than its depth.

Figure 20 is a view of the distal end of catheter 408 showing another 15 therapeutic device disclosed thereon. In Figure 20 an abrasive burr 418 is shown extending transversely from catheter 408. When rotated, burr 14 can form a crater 444. In both Figures 19 and 20, electrode 416 and burr 418 are shown spaced from heart wall 36. While creating craters 444, it is understood that loop 404 will be deployed in second position B such that electrode 416 and burr 418 20 will be in contact with heart wall 36.

It can be appreciated that each of the devices disclosed herein can be bi-polar as well as mono-polar. To make a bi-polar configuration, a ground

electrode would need to be disposed on the device proximate the electrode(s) shown.

Numerous characteristics and advantages of the invention covered by this document have been set forth in the foregoing description. It will be understood, 5 however, that this disclosure is, in many respects, only illustrative. Changes may be made in details, particularly in matters of shape, size, and arrangement of parts without exceeding the scope of the invention. The invention's scope is, of course, defined in the language in which the appended claims are expressed.

What is claimed is:

1. A method for increasing blood perfusion to heart muscle wall by forming a plurality of channels in the myocardium comprising the steps:
  - marking a first location within said heart muscle wall by securing a radiopaque marker to said heart muscle wall;
  - positioning a radiopaque, channel forming tip at a second location with a chamber of said heart;
  - viewing said first and second locations fluoroscopically;
  - adjusting said second location relative to said first position; and
  - forming said channel in said myocardium at said second location using said treatment tip.
2. A method for increasing blood perfusion as recited in claim 1, wherein said radiopaque marker is secured to said heart muscle with a marker selected from the group of fasteners consisting of hooks, barbs, bent members and curled members.
3. A method for increasing blood perfusion as recited in claim 1, wherein said radiopaque marker is secured to said heart muscle adhesively.
4. A method for increasing blood perfusion as recited in claim 1, wherein said radiopaque marker is secured to said heart muscle by injecting said marker within said wall.

5. A method for increasing blood perfusion as recited in claim 1, wherein said fluoroscopic visualization is accomplished on a computer generated display.

6. A method for increasing blood perfusion as recited in claim 4, wherein said radiopaque marker is a radiopaque dye.

7. A method for increasing blood perfusion as recited in claim 4, wherein said radiopaque marker is a metal.

8. A method for increasing blood availability to heart muscle myocardium having healthy, hibernating, and infarcted dead tissue comprising:  
providing a treatment tip having the capability of forming a channel in said myocardium from within said heart;  
selecting a first location in said healthy tissue;  
selecting a second location in said hibernating tissue; and  
forming a plurality of channels utilizing said treatment tip in said myocardium in a pattern extending from said first location to said second location.

9. A method as recited in claim 8, wherein said channels are formed substantially sequentially.

10. A method as recited in claim 8, wherein said treatment tip includes a plurality of cutting tips and said channels are formed substantially at the same time.

11. A method as recited in claim 8, wherein said pattern is substantially linear between said first and second locations.

12. A method as recited in claim 8, wherein said pattern is substantially a circular cluster including said first and second locations.

13. A method as recited in claim 8, wherein said pattern is substantially an array between said first and second locations.

14. A method as recited in claim 8, wherein said treatment tip is selected from the group consisting of mechanical cutting probes, laser cutting probes and radio frequency cutting probes.

15. A method as recited in claim 8, further comprising injecting radiopaque material into said channels, such that the positions of said channels are fluoroscopically viewable.

16. A method as recited in claim 8, wherein said treatment tip includes a mechanical cutting tip having a lumen therethrough, further comprising

injecting radiopaque material through said lumen into said channels, such that the position of said channels are fluoroscopically viewable.

17. A method for increasing blood availability to heart muscle by forming a plurality of channels in the myocardium comprising the steps:

providing a myocardial channel forming device having an anchoring member, at least one treatment member, and means for rotating said treatment member about said anchoring member, wherein said treatment member includes means for forming a channel in said myocardium;

attaching said anchoring member to a first position in said myocardium;

rotating said treatment member to a desired rotational angle relative to said anchor; and

forming a channel in said myocardium.

18. A method for increasing blood availability to heart muscle as recited in claim 17, wherein

said channel forming device has a longitudinal axis extending through said anchoring member,

said treatment member has a distal tip displaced a radial distance from said longitudinal axis,

said channel forming device includes means for controlling said treatment member distal tip radial distance,

further comprising setting said treatment member radial distance.

19. A method for increasing blood availability to heart muscle as recited in claim 18, wherein said means for controlling said rotational angle includes an elongate tubular member having a lumen and a wall, wherein said anchoring member is disposed within said lumen and said treatment probe is attached to said tube wall, such that said treatment member rotational angle can be controlled by rotating said elongate tube about said anchoring member.

20. A method for increasing blood availability to heart muscle as recited in claim 19, wherein said treatment member is slidably attached to said tubular wall, said treatment member distal tip is bent away from said anchoring member and has a longitudinal displacement relative to said anchoring member, such that the longitudinal and radial displacement of said treatment member distal tip can be adjusted by slidably adjusting said treatment member distal tip longitudinal displacement within said tube.

21. A method for increasing blood availability to heart muscle as recited in claim 18, wherein said device includes an elongate tubular member having a lumen and a wall, wherein said anchoring member and said treatment probe are disposed within said lumen, such that said treatment member rotational angle can be controlled by rotating said treatment member.

22. A device for creating a plurality of myocardial channels comprising:

an elongate tube having a distal port;

an anchoring member slidably disposed within said elongate tube, said anchoring member having a pointed distal end extending from said elongate tube distal port, said pointed distal end adapted to penetrate into said myocardium;

a treatment member slidably disposed within said elongate tube, said treatment member having an arcuate distal region having a cutting tip thereon, said arcuate distal region being biased to increase in radius when not constrained within said tube, such that said distally extending said elongate tube causes said treatment member distal end to decrease in radial distance from said anchor member and proximally retracting said elongate tube causes said treatment member distal end to increase in radial distance from said anchor member.

23. A device for creating a plurality of myocardial channels as recited in claim 22, wherein said treatment member is disposed within a treatment member lumen in said elongate tube distinct from said anchoring member lumen, such that said treatment member can be rotated about said anchoring member by rotating said elongate tube about said anchoring member.

24. A device for creating a plurality of myocardial channels as recited in claim 23, further comprising means for restricting said rotation of said treatment member relative to said anchoring member to a finite set of rotational positions.

25. A device for creating a plurality of myocardial channels comprising:

an elongate tube having a distal port and a longitudinal axis;  
a plurality of wires disposed within said elongate tube, said wires having distal tips deployed about said longitudinal axis, such that a pattern of wire distal tips is formed distal of said tube distal port;  
a radio-frequency energy source connected to said wires, said energy source capable of creating energy at said wire distal tips sufficient to burn channels in said myocardium, such that a pattern of myocardial channels can be created by supplying said radio-frequency energy to said wires.

26. A method as recited in claim 8, further providing at least one external magnet, wherein said treatment tip has a distal region responsive to magnetic forces, such that said treatment tip is at least partially guided into a cutting position utilizing said magnetically responsive distal region and said external magnet.

27. A device for creating a myocardial channel comprising:  
a catheter having a magnetically responsive distal region and a distal cutting tip, such that said cutting tip can be at least partially guided into a cutting position utilizing said magnetically responsive distal region responding to an external magnetic force.

28. A device for creating a myocardial channel as recited in claim 27, wherein said catheter magnetically responsive distal region can be pulled by said external magnetic force.

29. A device for lessening the movement of a beating heart wall portion comprising:

a catheter having a magnetically responsive distal region, said catheter being adapted to be received within a coronary artery, such that said catheter magnetically responsive distal region can have force exerted thereon by an external magnet while said catheter is received within said coronary artery.

30. A method for lessening the movement of a beating heart wall portion comprising the steps:

providing a catheter having a magnetically responsive distal region;  
providing at least one external magnet;  
inserting said catheter distal region into a coronary vessel near said heart wall portion; and  
positioning said external magnet near said catheter distal region, such that said external magnet exerts force on said magnetically responsive catheter distal region within said heart wall portion, presenting resistance to said heart wall movement.

31. A device for lessening the movement of a beating heart wall portion comprising:

a catheter having a magnetically responsive distal region, said catheter being adapted to be received within a coronary artery, such that said catheter magnetically responsive distal region can have force exerted thereon by an external magnet while said catheter is received within said coronary artery.

32. A device for lessening said beating heart wall portion movement as recited in claim 31, wherein said magnetically responsive distal region can be pulled by said external magnet.

33. A method for increasing blood availability to heart muscle as recited in claim 17, further providing at least one external magnet, wherein said anchoring member includes a magnetically responsive distal region and said attaching step includes exerting magnetic force by said external magnet on said anchor member magnetically responsive distal region, such that said anchor member distal region is forced toward said myocardium.

34. A device for creating a plurality of myocardial channels comprising:

an elongate tube having a distal port;  
an anchoring member slidably disposed within said elongate tube, said anchoring member having a magnetically responsive distal region extending from

said elongate tube distal port, said distal region adapted to contact a heart wall region near said myocardium, such that said anchoring member can be forced into said heart wall region in response to an external magnetic force; and

a treatment member slidably disposed within said elongate tube, said treatment member having an arcuate distal region having a cutting tip thereon, said arcuate distal region being biased to increase in radius when not constrained within said tube, such that said distally extending said elongate tube causes said treatment member distal end to decrease in radial distance from said anchor member and proximally retracting said elongate tube causes said treatment member distal end to increase in radial distance from said anchor member.

35. A device for creating a plurality of myocardial channels as recited in claim 34, wherein said treatment member is disposed within a treatment member lumen in said elongate tube distinct from said anchoring member lumen, such that said treatment member can be rotated about said anchoring member by rotating said elongate tube about said anchoring member.

36. A device for guiding a PMR cutting wire distal region to a plurality of heart chamber wall sites comprising:

an elongate outer tube having a tubular wall and a distal region, said distal region wall having a plurality of channels therethrough; and  
means for stabilizing the position of said outer tube distal region within said heart chamber,

such that said PMR cutting wire distal region can be advanced through said outer tube distal region wall channels to said heart chamber wall sites.

37. A device for guiding a PMR cutting wire as recited in claim 36 wherein said means for stabilizing said outer tube position includes a suction orifice disposed in said outer tube distal region, said orifice being adapted to make contact with said heart chamber wall, said orifice being in fluid communication with a vacuum lumen substantially co-extensive with said outer tube.

38. A device for guiding a PMR cutting wire as recited in claim 36 wherein said means for stabilizing said outer tube position includes a magnetically responsive portion of said outer tube distal region, such that an externally applied magnetic field can exert a force upon said outer tube distal region.

39. A device for creating a plurality of myocardial channels in a heart chamber wall comprising:

an elongate outer tube having a tubular wall and a distal region, said distal region wall having a plurality of channels therethrough;  
means for stabilizing the position of said outer tube distal region within said heart chamber; and

a PMR cutting wire having a distal region disposed within said outer tube,

such that said PMR cutting wire distal region can be advanced through said outer tube distal region wall channels to said heart chamber wall.

40. A device as recited in claim 39, further comprising a guide tube disposed about said PMR cutting wire and disposed within said outer tube.

41. A device as recited in claim 39, wherein said PMR cutting wire distal region has a distal tip and an arcuate bias proximate said distal tip such that said distal region is predisposed to extend through said outer tube channels when said distal tip is passed over said channels.

42. A device as recited in claim 39, wherein said means for stabilizing said outer tube position includes a suction orifice disposed in said outer tube distal region, said orifice being adapted to make contact with said heart chamber wall, said orifice being in fluid communication with a vacuum lumen substantially co-extensive with said outer tube.

43. A device as recited in claim 39, wherein said means for stabilizing said outer tube position includes a magnetically responsive portion of said outer tube distal region, such that an externally applied magnetic field can exert a force upon said outer tube distal region.

44. A device as recited in claim 39, further comprising an elongate shape member said outer tube and secured to said outer tube wall over at least said outer tube distal region, said shape member having a preformed shape such that said outer tube distal region is biased to assume said shape member preformed shape.

45. A device for creating a plurality of myocardial channels in a heart chamber wall comprising:

an elongate outer tube having a longitudinal axis, a proximal end, a distal end, a tubular wall and a distal region, said distal region wall having an elongate slot therethrough, said slot having a length;

an elongate intermediate tube disposed within said outer tube, said intermediate tube having a wall, a distal region, and a channel in said distal region wall; and

an elongate inner PMR cutting probe disposed within said intermediate tube and adapted to extend through said intermediate tube channel such that a distal length of said PMR cutting probe extends outside of said intermediate and outer tubes, wherein said PMR cutting probe extends at a longitudinal position relative to said outer tube,

said outer tube slot having a proximal portion and a distal portion, said intermediate tube being slidable within said outer tube, said inner PMR cutting probe being slidable within said intermediate tube,

such that said PMR probe extended distal length can be varied by sliding said inner PMR probe within said intermediate tube and through said intermediate tube channel,

such that said PMR probe longitudinal position can be varied by sliding said intermediate tube within said slot,

such that said PMR probe can be rotated by rotating said outer tube.

46. A device for creating a plurality of myocardial channels in a heart chamber wall comprising:

an elongate rod having a proximal region and a distal region, said rod distal region secured to an outer collar;

an intermediate tube slidably received within said collar;

an inner PMR cutting probe slidably received within said intermediate tube;

an elongate anchoring member having a proximal region and a distal end, said anchoring member being slidably secured to said collar, said anchoring member having anchoring means disposed proximate said anchoring member distal end,

said intermediate tube distal region containing said inner PMR probe together having a distal region arcuate bias, such that extending said outer collar distally over said intermediate tube straightens out said intermediate tube and retracting said outer collar proximally over said intermediate tube allows said arcuate shape to return,

such that sliding said intermediate tube relative to said anchoring member moves said contained inner PMR probe distal end relative to said anchoring means.

47. A device as recited in claim 46, wherein said anchoring means includes a pigtail adapted to screw into said heart chamber wall and said anchoring member is rotatable relative to said collar, such that said pigtail can be rotated by rotating said anchoring member.

48. A device as recited in claim 46, wherein said anchoring means includes a magnetically responsive portion, such that an externally applied magnetic field can exert a force upon said anchoring member distal region.

49. A method for increasing blood availability to heart muscle by causing localized tissue death at a plurality of sites in the myocardium comprising the steps:

providing a myocardial localized tissue killing device having an anchoring member, at least one treatment member, and means for rotating said treatment member about said anchoring member, wherein said treatment member includes means for killing tissue in said myocardium;

attaching said anchoring member to a first position in said myocardium;

rotating said treatment member to a desired rotational angle relative to said anchor; and

causing tissue death in said myocardium.

50. A method as recited in claim 49, wherein said treatment member includes a tube having a lumen therethrough and said causing step includes delivering a cold substance through said lumen.

51. A method as recited in claim 50, wherein said cold substance is liquid nitrogen.

52. A method as recited in claim 50, wherein said cold treatment member has a distal orifice in fluid communication with said treatment member lumen.

53. A device for killing tissue at a plurality of myocardial sites comprising:

an elongate tube having a distal port;  
an anchoring member slidably disposed within said elongate tube, said anchoring member having a pointed distal end extending from said elongate tube distal port, said pointed distal end adapted to penetrate into said myocardium;  
a treatment member capable of causing localized tissue death slidably disposed within said elongate tube, said treatment member having an arcuate distal region having a treatment tip thereon, said arcuate distal region being biased to increase in radius when not constrained within said tube, such that said distally

extending said elongate tube causes said treatment member distal end to decrease in radial distance from said anchor member and proximally retracting said elongate tube causes said treatment member distal end to increase in radial distance from said anchor member, wherein said treatment member is disposed within a treatment member lumen in said elongate tube distinct from said anchoring member lumen, such that said treatment member can be rotated about said anchoring member by rotating said elongate tube about said anchoring member.

54. A device as recited in claim 53, wherein said treatment member includes a lumen adapted to deliver liquid nitrogen to said treatment member distal end.

55. A device as recited in claim 54, wherein said treatment member distal end includes an orifice in communication with said treatment member lumen.

56. A device for causing localized tissue death at a plurality of myocardial sites comprising:

an elongate tube having a distal port and a longitudinal axis;  
a plurality of treatment tubes disposed within said elongate tube, said treatment tubes having distal tips deployed about said longitudinal axis, such that a pattern of treatment tube distal tips is formed distal of said tube distal port;

a cryogenic substance source connected to said treatment tubes, said cryogenic source capable of causing cold at said treatment tube distal tips sufficient to cause tissue death in said myocardium, such that a pattern of localized myocardial tissue death can be created by supplying said cryogenic substance to said treatment tubes.

57. A device as recited in claim 56, wherein said cryogenic substance is liquid nitrogen.

58. A catheter assembly, comprising:

a guidewire having a proximal end and a distal end;

an expandable member disposed at the distal end of the guide wire, the expandable member being moveable between a first position and a second position, in the first position, the member being collapsed to move through a lumen of a guide catheter, in a second position, the expandable member having a transverse diameter with respect to the length of the guidewire greater than the transverse diameter of the guide catheter lumen;

an elongate catheter having a proximal end and a distal end, the catheter defining an elongate lumen, and the guidewire being disposed in the lumen; and

a therapeutic device connected to the catheter and disposed proximate the distal end of the catheter.

59. The catheter assembly in accordance with claim 58, wherein the expandable member includes a wire loop.

60. The catheter assembly in accordance with claim 58, wherein the therapeutic device includes a needle.

61. The catheter assembly in accordance with claim 60, wherein the needle includes a hypotube.

62. The catheter assembly in accordance with claim 58, wherein the therapeutic device includes an electrode.

63. The catheter assembly in accordance with claim 62, wherein the electrode extends transversely from the catheter a length greater than width of the electrode.

64. The catheter assembly in accordance with claim 58, wherein the therapeutic device includes a rotatable burr.

65. A method of performing PMR, comprising the steps of:  
providing a catheter assembly including a guidewire having a distal end expandable member, and a catheter advancable over the guidewire, the catheter having a distal end and a therapeutic device disposed proximate the distal end;

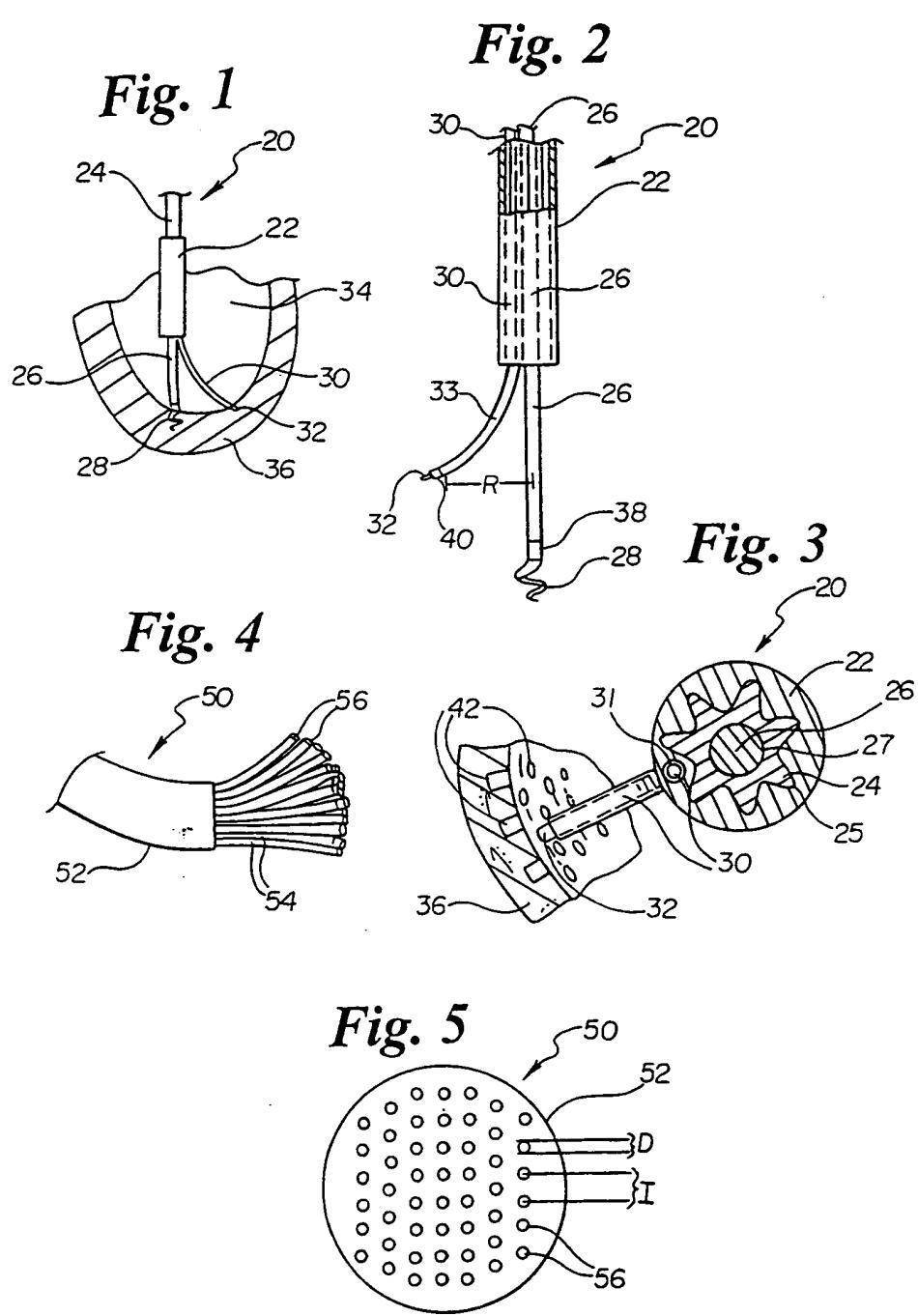
advancing the expandable member into a chamber of a patient's heart;  
expanding the expandable member such that a length of the guidewire  
proximate the distal end of the guidewire is disposed proximate a wall of the  
chamber;

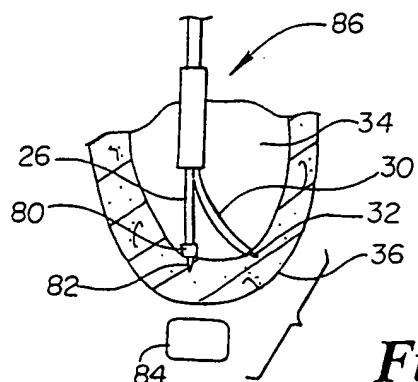
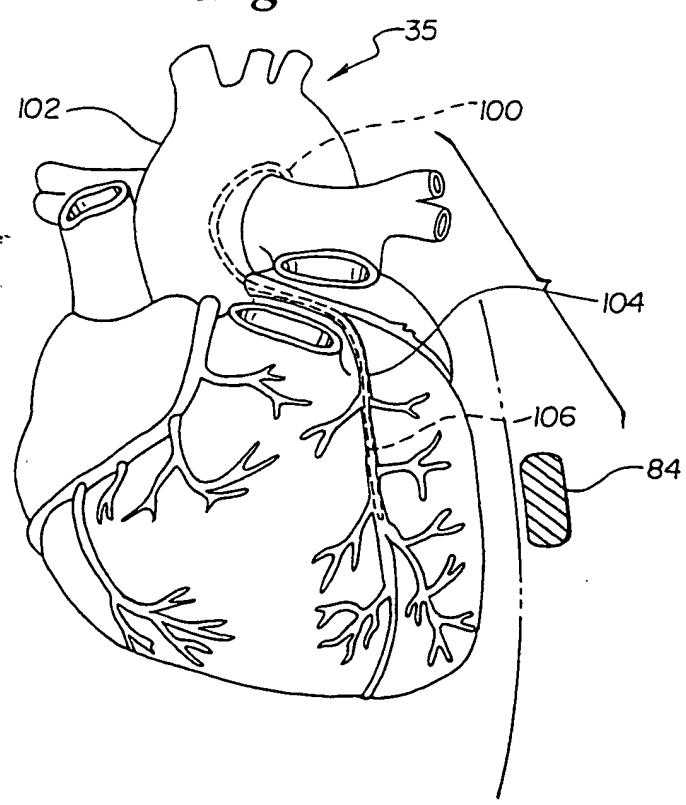
advancing the catheter over the guidewire such that the therapeutic device  
is adjacent the wall; and

actuating the therapeutic device to deliver therapy to the heart wall.

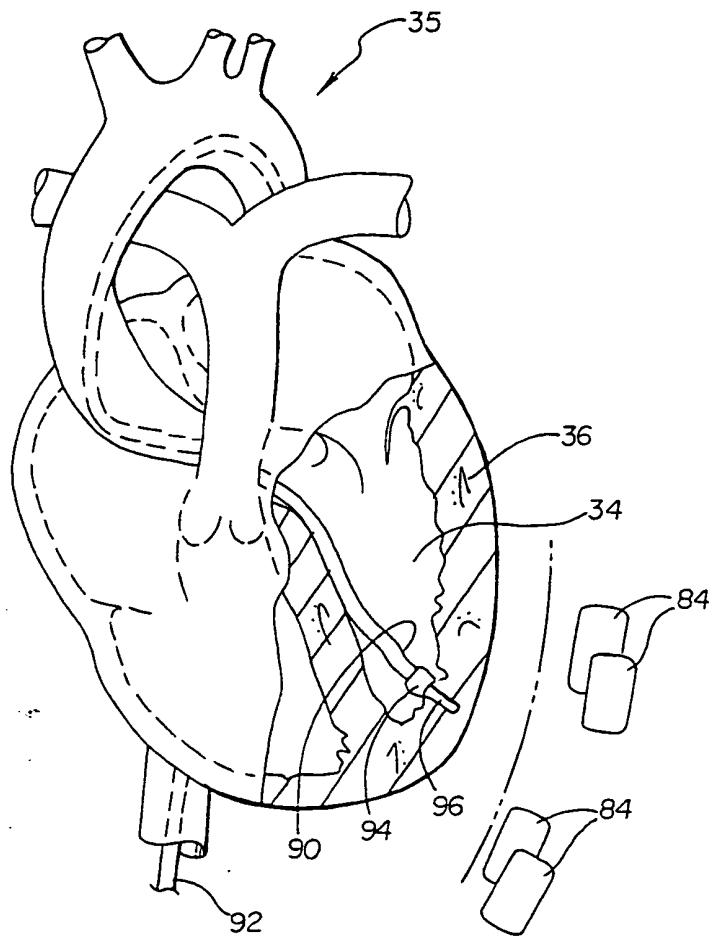
66. The method in accordance with claim 65, further comprising the  
step of incrementally advancing the therapeutic device along the heart wall and  
delivering the therapy to the heart wall intermittently as the device is  
incrementally advanced.

67. The method in accordance with claim 65, further comprising the  
step of rotating the guidewire to disposed the catheter proximate another portion  
of the wall.

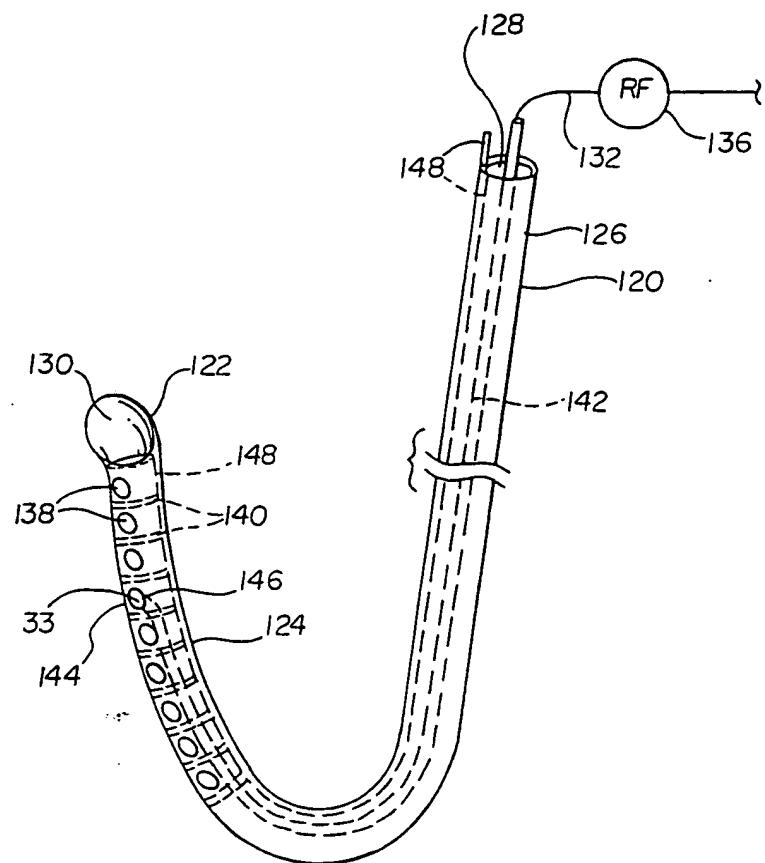


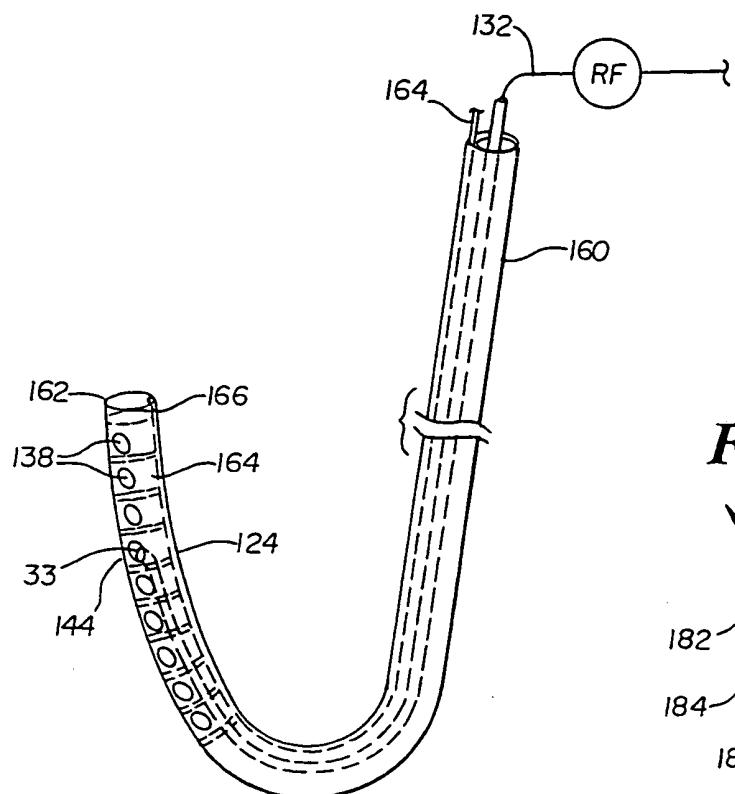
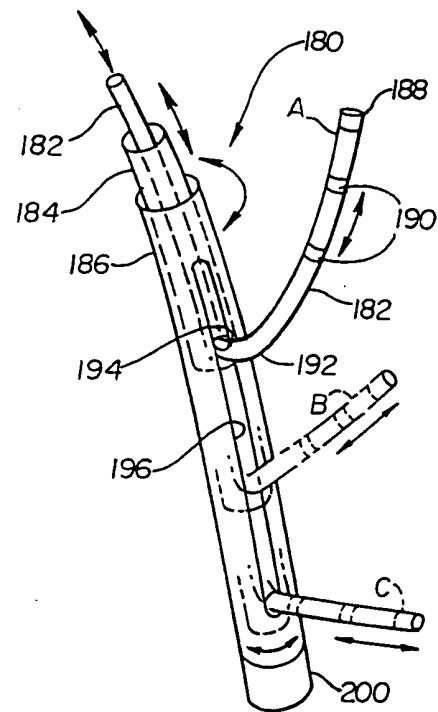
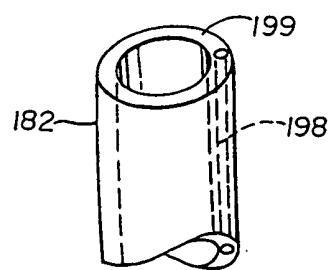
*Fig. 6**Fig. 8*

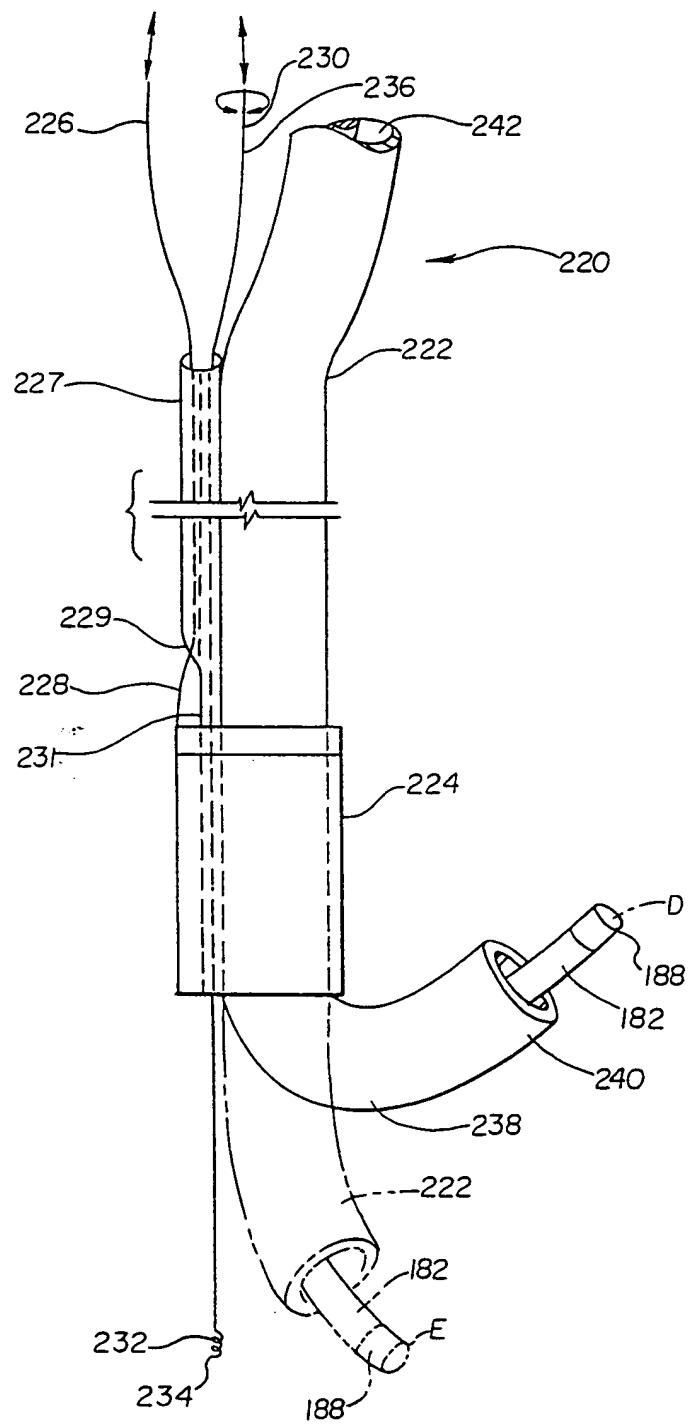
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*Fig. 7*

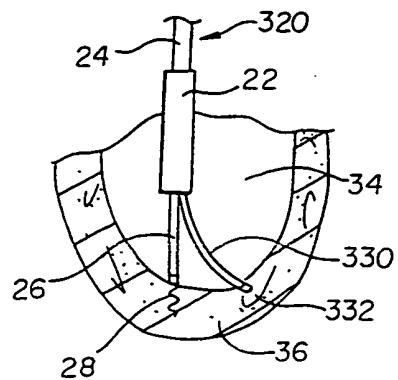
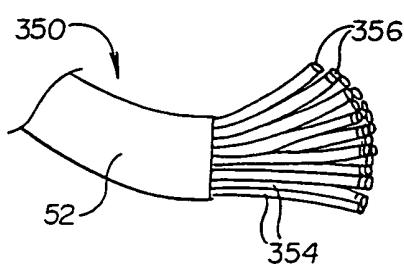
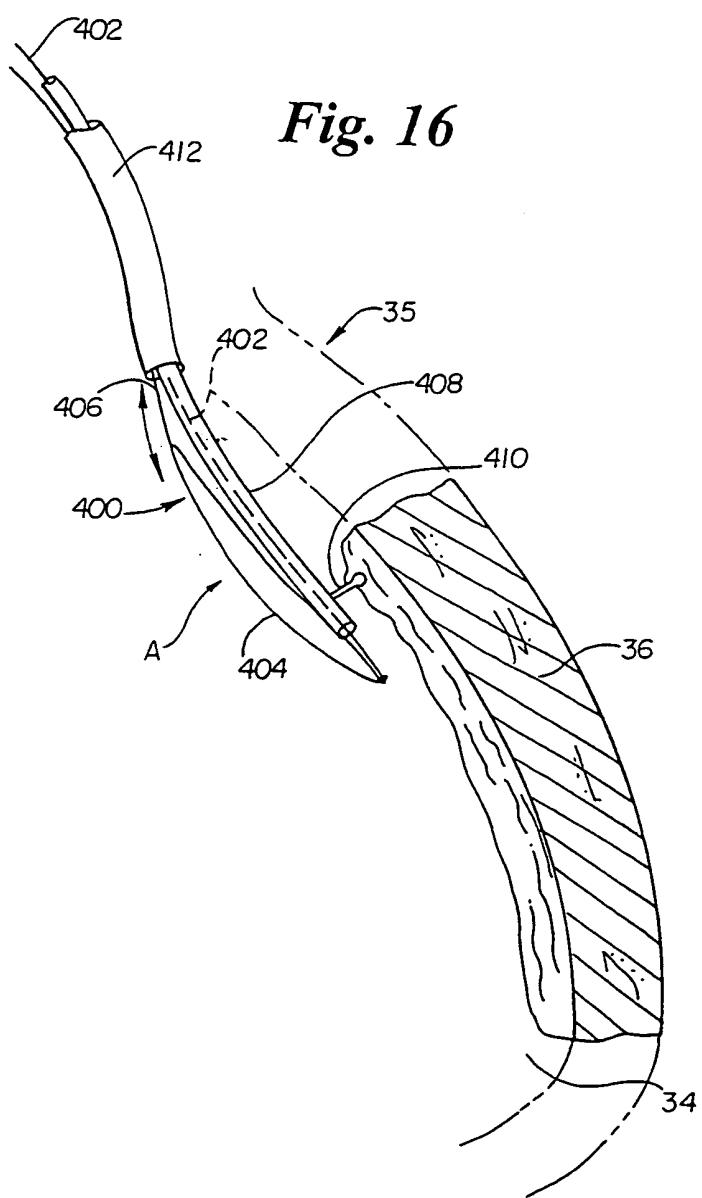
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*Fig. 9*

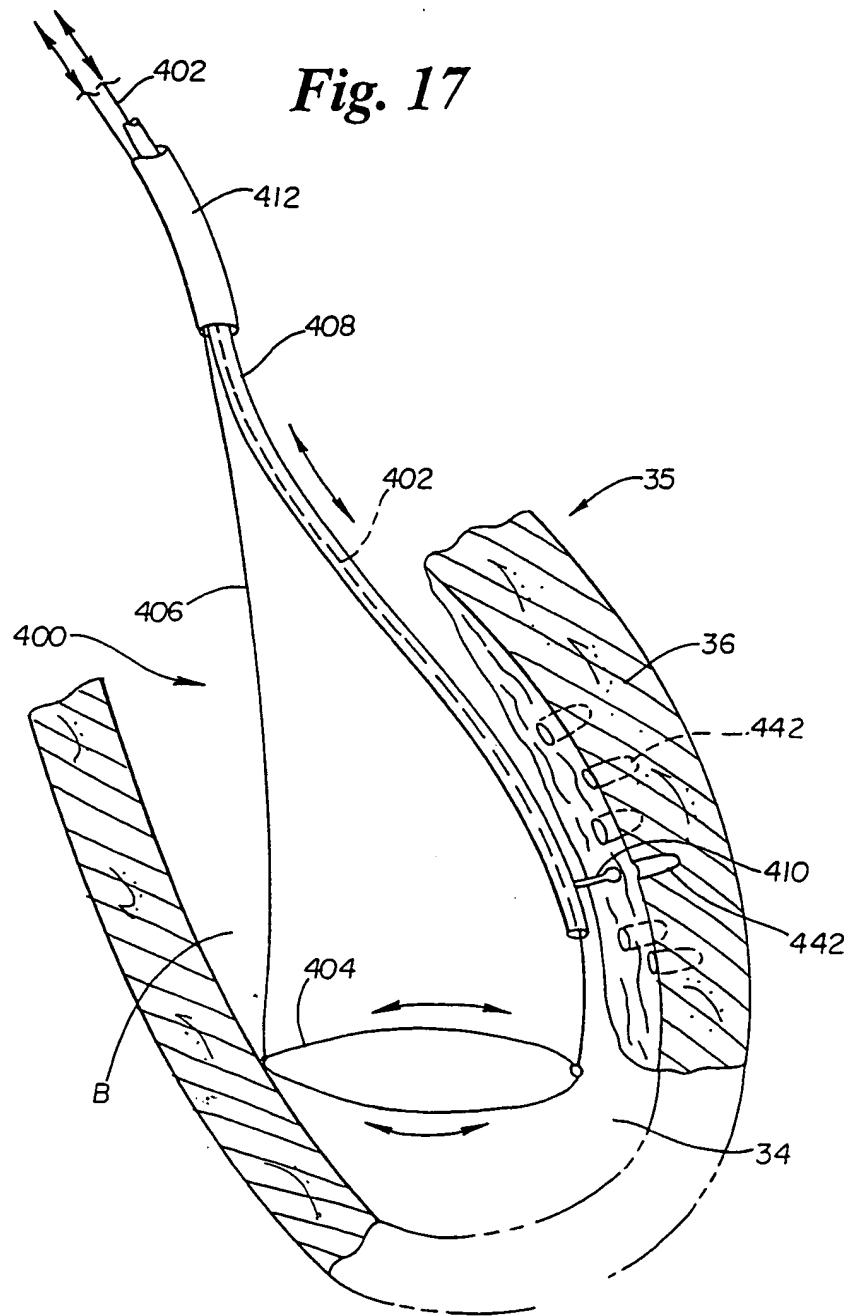
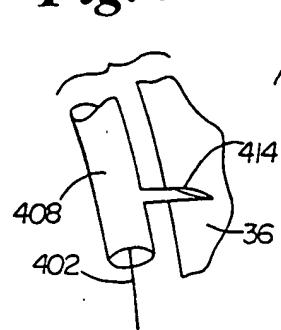
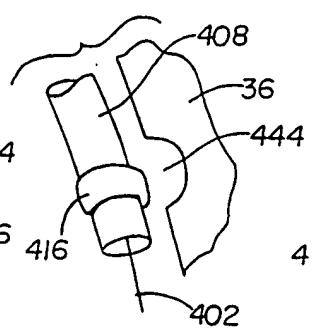
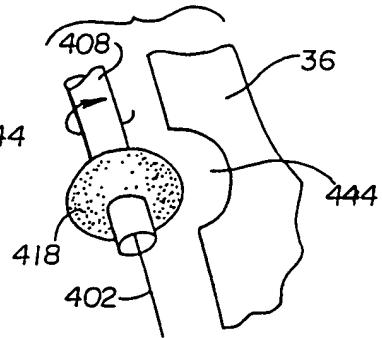
*Fig. 10**Fig. 11**Fig. 12*

*Fig. 13*

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*Fig. 14**Fig. 15**Fig. 16*

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*Fig. 17**Fig. 18**Fig. 19**Fig. 20*

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US98/23397

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) : A61B 17/36

US CL : 606/15

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 606/8-13, 15-17

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,593,405 A (OSYPKA) 14 January 1997.	22-24, 34, 35, 52-57
A, P	US 5,703,985 A (OWYANG) 30 December 1997.	22-24, 34, 35, 53-57
Y, E	US 5,840,075 A (MUELLER et al) 24 November 1998.	22-24, 34, 35, 53-57
Y,P	US 5,782,823 A (MUELLER) 21 July 1998.	22-24, 34, 35, 53-57
A	US 5,423,804 A (KULICK) 13 June 1995.	22-24, 34, 35, 53-57
A	US 5,437,660 A (JOHNSON et al) 01 August 1995.	22-24, 34, 35, 53-57

Further documents are listed in the continuation of Box C.  See patent family annex.

* Special categories of cited documents:	"T"	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"A" document defining the general state of the art which is not considered to be of particular relevance	"X"	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y"	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&"	document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed		

Date of the actual completion of the international search

20 JANUARY 1999

Date of mailing of the international search report

03 MAR 1999

Name and mailing address of the ISA/US  
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## INTERNATIONAL SEARCH REPORT

International application No.

PCT/US98/23397

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,498,258 A (HAKKY et al) 12 March 1996.	22-24, 34, 35, 53-57

**INTERNATIONAL SEARCH REPORT**

International application No.

PCT/US98/23397

**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1.  Claims Nos.: because they relate to subject matter not required to be searched by this Authority, namely:
  
2.  Claims Nos.: because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3.  Claims Nos.: because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

**Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)**

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1.  As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2.  As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3.  As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4.  No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: 22-24, 34, 35, 53-57

**Remark on Protest**  

The additional search fees were accompanied by the applicant's protest.

No protest accompanied the payment of additional search fees.

**BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING**

This ISA found multiple inventions as follows:

Claims 1-7, method for increasing blood perfusion to the heart including radiopaque marking.

Claims 8-21, method for increasing blood availability.

Claims 25 and 26, a device for creating a plurality of channels having wires, and an RF source.

Claims 27-33, a magnetic catheter.

Claims 36-48 and 58-67, a device for guiding a PMR cutting wire.

Claims 22-24, 34, 35 and 53-57, a device for increasing blood availability.

Claims 49-52, a method for increasing blood availability.

Group I is distinct from Groups II-VII in that it is a method for increasing blood perfusion to the heart via use of radiopaque marking.

Group II is distinct from Groups I and III-VII in that it is a method for increasing blood availability via use of a treatment tip to make a plurality of channels.

Group III is distinct from Groups I, II and IV-VII in that it is a device for creating myocardial channels using a radio frequency energy source.

Group IV is distinct from Groups I-III and V-VII in that it is a device for creating a channel comprising a catheter having a magnetically responsive distal region.

Group V is distinct from Groups I-IV, VI and VII in that it is a device for guiding a cutting device including a tube, and stabilization means.

Group VI is distinct from Groups I-V and VII in that it is a device for creating a plurality of myocardial channels including the combination of an elongate tube, anchoring member, and a treatment member.

Group VII is distinct from Groups I-VI in that it is a method for increasing blood availability providing a localized tissue killing device with an anchoring member.

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